

## HUMAN ENDOCRINE- GLAND-DERIVED VASCULAR ENDOTHELIAL GROWTH FACTOR (EG-VEGF) ELISA KIT

FOR THE QUANTITATIVE DETERMINATION  
OF HUMAN EG-VEGF CONCENTRATIONS IN  
SERUM, EDTA PLASMA AND CELL  
CULTURES.



### PURCHASE INFORMATION:

ELISA Name	Human EG-VEGF ELISA
Catalog No.	SK00151-01
Formulation	96 T
Standard range	15.6-1000 pg/mL
Sensitivity	7.8 pg/mL
Sample Volume	100 µl
Sample Type	Serum, EDTA plasma, cell cultures
Dilution Factor	N/A
Specificity	Human EG-VEGF only
Calibration	Human EG-VEGF rec
Intra-assay Precision	6-8%
Inter-assay Precision	10-12%
Storage	2 °C-8 °C

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FOR RESEARCH USE ONLY. NOT FOR USE IN  
DIAGNOSTIC PROCEDURES.

## INTRODUCTION

Human EG-VEGF immunoassay is a 3.5 - 4.5 hour solid phase ELISA designed to measure human EG-VEGF in serum, and plasma. It contains recombinant human EG-VEGF and antibodies raised against this protein. It has been shown to accurately quantitate recombinant human EG-VEGF. Results obtained with naturally occurring EG-VEGF samples showed linear curves that were parallel to the standard curves obtained using the kit standards. These results indicate that the Immunoassay kit can be used to determine relative mass values for natural human EG-VEGF.

## PRINCIPLE OF THE ASSAY

This assay employs the quantitative sandwich enzyme immunoassay technique. A monoclonal antibody specific for EG-VEGF has been pre-coated onto a microplate. Standards and samples are pipetted into the wells and any EG-VEGF present is bound by the immobilized antibody. After washing away any unbound substances, a biotinylated polyclonal antibody specific for EG-VEGF is added to the wells. Following a wash to remove any unbound antibody-biotin reagent, a Streptavidin HRP link is added to the wells. After washing away any unbound enzyme, a substrate solution is added to the wells and color develops in proportion to the amount of EG-VEGF bound in the initial step. The color development is stopped and the intensity of the color is measured.

## LIMITATIONS OF THE PROCEDURE

- \_ FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES.
- \_ The kit should not be used beyond the expiration date on the kit label.
- \_ Do not mix or substitute reagents with those from other lots or sources.
- \_ It is important that the Dilution Buffer selected for the standard curve be consistent with the samples being assayed.
- \_ If samples generate values higher than the highest standard, dilute the samples with the appropriate Dilution Buffer and repeat the assay.
- \_ Any variation in standard diluent, operator, pipetting technique, washing technique, incubation time or temperature, and kit age can cause variation in binding.
- \_ This assay is designed to eliminate interference by soluble receptors, binding proteins, and other

factors present in biological samples. Until all factors have been tested in the Immunoassay, the possibility of interference cannot be excluded.

## MATERIALS PROVIDED

Description	Code	Quantity
<b>Microplate</b> - 96 well polystyrene microplate (12 strips of 8 wells) coated with a monoclonal antibody against EG-VEGF.	151-01-01	1 plate
<b>EG-VEGF Standard</b> – 1 ng/vial of recombinant human EG-VEGF in a buffered protein base with preservatives; lyophilized.	151-01-02	1 vial
<b>Detection Antibody Concentrate</b> – 120 µL / vial, 100-fold concentrated of Biotinylated polyclonal antibody against EG-VEGF with preservatives; lyophilized.	151-01-03	1 vial
<b>Positive Control</b> - one of recombinant human EG-VEGF, lyophilized	151-01-04	1 vial
<b>Streptavidin-HRP Conjugate</b> -120 ul/vial, 100-fold concentrated solution of Streptavidin conjugate to HRP	SHHRP	1 vial
<b>Dilution Buffer</b> - 60mL/vial of buffered protein based solution with preservatives	DB01	1 vial
<b>Wash Buffer</b> -50 ml/vial, 10-fold concentrated buffered surfactant, with preservative.	WB01	1 vial
<b>TMB Substrate Solution</b> -13 ml / vial of TMB substrate solution	TMB01	1 vial
<b>Stop Solution</b> (0.5M HCl) , 13 ml /vial of 0.5M HCl	S-STOP	1 vial
<b>Plate Covers</b> – Plate sealer.	EAPS	1

## STORAGE

**Unopened Kit:** Store at 2 - 8° C. Do not use past kit expiration date.

**Opened / Reconstituted Reagents:** May be stored for up to 1 month at 2 - 8°C.

**Standard :** Reconstituted standard should be stored for up to two weeks at -70° C. *Diluted standard working solution and Positive Control should be prepared and used immediately.*

**Microplate Wells:** Return unused wells to the plastic bag containing the desiccant pack, reseal along entire edge of zip-seal. May be stored for up to 8 months at 2 - 8° C.

**OTHER SUPPLIES REQUIRED**

- Microplate reader capable of measuring absorbance at 450 nm, with the correction wavelength set at 540 nm or 570 nm.
- Microplate shaker (250-300rpm).
- Pipettes and pipette tips.
- Deionized or distilled water.
- Squirt bottle, manifold dispenser, or automated microplate washer.
- 100 mL and 500 mL graduated cylinders.

**SAMPLE COLLECTION AND STORAGE**

**Cell Culture Supernates** - Remove particulates by centrifugation and assay immediately or aliquot and store samples at ≤-20° C. Avoid repeated freeze-thaw cycles.

**Serum** - Use a serum separator tube (SST) and allow samples to clot for 30 minutes before centrifugation for 15 minutes at 1000 x g. Remove serum and assay immediately or aliquot and store samples at ≤ -20° C. Avoid repeated freeze-thaw cycles.

**Plasma** - Collect plasma using EDTA, heparin, or citrate as an anticoagulant. Centrifuge for 15 minutes at 1000 x g within 30 minutes of collection. Assay immediately or aliquot and store samples at ≤-20° C. Avoid repeated freeze-thaw cycles.

**PRECAUTIONS FOR USE**

All reagents should be considered as potentially hazardous. The stop solution contains diluted Hydrochloric acid. Appropriate care, therefore, should be taken while handling this solution. We therefore recommend that this product is handled only by those persons who have been trained in laboratory techniques and that it is used in accordance with the principles of good laboratory practice. Wear suitable protective clothing such as laboratory overalls, safety glasses and gloves. Care should be taken to avoid contact with skin or eyes. In

the case of contact with skin or eyes wash immediately with water.

**REAGENT PREPARATION**

**Bring all reagents to room temperature before use.**

**Wash Buffer** - If crystals have formed in the concentrate, warm to room temperature and mix gently until the crystals have completely dissolved. Dilute 50 mL of Wash Buffer Concentrate into deionized or distilled water (450 mL) to prepare 500 mL of Wash Buffer.

**EG-VEGF Standard - Refer to vial label for reconstitution volume.** Reconstitute the **EG-VEGF Standard** with 1 ml of Dilution Buffer. This reconstitution produces a stock solution of 1000 pg/mL. Allow the standard to sit for a minimum of 15 minutes with gentle agitation prior to making dilutions. Pipette 250 µL of the appropriate Dilution Buffer into the tube #1 and 6. Use the stock solution to produce a dilution series (below). Mix each tube thoroughly before the next transfer. The 1000 pg/mL standard serves as the high standard. The appropriate Dilution Buffer serves as the zero standard (0 pg/mL).

Standard	Standard	Reagent Diluent	Concentration
stock	Powder	1000 µl	1000 pg/ml
# 1	100 µl of stock	900 µl	500 pg/ml
# 2	250 µl of 1	250 µl	250 pg/ml
# 3	250 µl of 2	250 µl	125 pg/ml
# 4	250 µl of 3	250 µl	62.5 pg/ml
# 5	250 µl of 4	250 µl	31.25 pg/ml
# 6	250 µl of 5	250 µl	15.6 pg/ml

**Detection Antibody-** Reconstitute the **Detection Antibody concentrated** with 120 µl of Dilution Buffer to produce a 100-fold concentrated stock solution. Pipette 11. 88 mL of the appropriate Dilution Buffer into the 15 ml centrifuge tube and transfer 120 µl of 100-fold concentrated stock solution to prepare working solution.

**Streptavidin-HRP Conjugate** - Pipette 11. 88 mL of Dilution Buffer into the 15 ml centrifuge tube and

transfer 120 µl of 100-fold concentrated stock solution to prepare working solution.

### ASSAY PROCEDURE

**Bring all reagents and samples to room temperature before use. It is recommended that standards be assayed in duplicate.**

1. Prepare all reagents and working standards as directed in the previous sections.
2. Remove excess micro-plate strips from the plate frame, return them to the foil pouch containing the desiccant pack, reseal.
3. Add 100 µL of Dilution Buffer to Blank well.
4. Add 100 µL of Standard, sample, or control per well. Cover with the Sealer. Incubate for 2 hours on micro-plate shaker at room temperature. A plate layout is provided to record standards and samples assayed.
5. Aspirate each well and wash, repeating the process three times for a total of four washes. Wash by filling each well with Wash Buffer (250 µL) using a squirt bottle, manifold dispenser, or autowasher. Complete removal of liquid at each step is essential to good performance. After the last wash, remove any remaining Wash Buffer by aspirating or decanting. Invert the plate and blot it against clean paper towels.
6. Add 100 µL of Detection Antibody working solution to each well. Cover with sealer. Incubate for 2 hours on micro-plate shaker at room temperature.
7. Repeat the aspiration/wash as in step 5.
8. Add 100 µL of **Streptavidin-HRP Conjugate** working solution to each well. Incubate for 40min on micro-plate shaker at room temperature.
9. Repeat the aspiration/wash as in step 5.
10. Add 100 µL of Substrate Solution to each well. Incubate for 20-30 minutes at room temperature. **Protect from light.**
11. Add 100 µL of Stop Solution to each well. The color in the wells should change from blue to yellow. If the color in the wells is green or if the color change does not appear uniform, gently tap the plate to ensure thorough mixing.
12. Determine the optical density of each well within 30 minutes, using a micro-plate reader set to 450 nm.

### CALCULATION OF RESULTS

Average the duplicate readings for each standard, control, and sample and subtract the average zero standard optical density. Create a standard curve

by reducing the data using computer software capable of generating a four parameter logistic (4-PL) curve fit. As an alternative, construct a standard curve by plotting the mean absorbance for each standard on the y-axis against the concentration on the x-axis and draw a best fit curve through the points on the graph. The data may be linearized by plotting the log of the EG-VEGF concentrations versus the log of the O.D. and the best fit line can be determined by regression analysis. This procedure will produce an adequate but less precise fit of the data. If samples have been diluted, the concentration read from the standard curve must be multiplied by the dilution factor.

### SUMMARY OF ASSAY PROCEDURE

PREPARE REAGENTS, SAMPLES AND STANDARDS
↓
Add 100µl of standard, samples, positive control to each well. Incubate 2 hours on the plate shaker at RT.
↓
Aspirate and wash 4 times.
↓
Add 100 µl Detection Antibody to each well. Incubate 2 hours on the plate shaker at RT.
↓
Aspirate and wash 4 times.
↓
Add 100 µl <b>Streptavidin-HRP Conjugate</b> to each well. Incubate 40 min on the plate shaker at RT.
↓
Aspirate and wash 4 times.
↓
Add 100 µl Substrate to each well. Incubate 20-30min on the bench top. Protect from light.
↓
Add 100 µl Stop Solution to each well. Read 450nm within 15min

### SPECIFICITY

PROTEIN	CROSSREACTIVITY
Human EG-VEGF	100%
Mouse EG-VEGF	0
Human VEGF-D	0

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Human VEGF165	0
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